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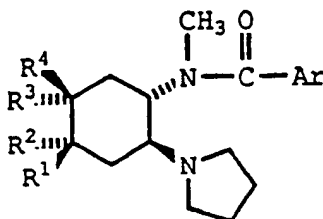
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(54) **1,2-cyclohexylaminoaryl amides useful as analgesic agents.**

(57) Novel 1,2-cyclohexylaminoaryl amides of formula I



I

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are useful as analgesic agents having very high kappa-opioid affinity and selectivity and potency and are useful as analgesics, diuretics, antiinflammatory and psychotherapeutic agents. Methods for making the compounds and pharmaceutical compositions containing them are also disclosed.

The search for strong analgesics which also possess minimal potential for dependency has been among the highest priority efforts in pharmacological research. These research efforts have, to a great extent, involved chemical modifications of the opiate structure and the discovery of chemically novel compounds which possess morphine-like activity.

The concept of multiple opioid receptors has been supported by studies with nalorphine and a series of benzomorphans which display unusual pharmacological properties dissimilar from morphine, yet blocked by the opioid antagonists. [See, for example, W. R. Martin, et al., J. Pharmacol. Exp. Ther., 197:517-531 (1976)].

The existence of multiple types of opioid receptors is of importance because it suggests the possibility of separating the desirable analgesic and psychotherapeutic effects of a drug compound from the undesirable abuse potential or habituating effect.

15 United States Patent 4,145,435 describes certain 2-amino-cycloaliphatic amide compounds as analgesic. In particular, trans-3,4-dichloro-N-methyl-N-[2-(1-pyrrolidiny)cyclohexyl]benzeneacetamide has been reported to possess selective kappa opioid agonist activity, and therefore to possess analgesic activity without attendant dependence liability. [See P. V. Vonvoigtlander, et al., J. Pharmacol. Exp. Ther., 224:7-12 (1983)].

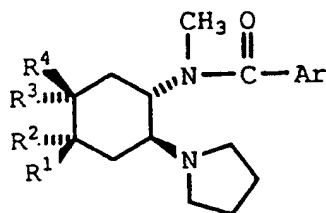
20 Recently, the diuretic effect of various opioid agonists and antagonists has been studied, and it has been shown that kappa agonists tend to increase urination, while mu agonists decreased urination. [See J. D. Leander, J. Pharmacol. Exp. Ther., 227:35-41 (1983)]. These findings suggest that opioid agonists and antagonists also possess potential as diuretics.

United States Patent 4,656,182 describes certain trans-1,2-diaminocyclohexyl amides useful as anal-
25 gesics, diuretics, and psychotherapeutics.

United States Patent 4,463,013 discloses certain oxygen substituted amino-cyclohexyl-benzeneacetamides as diuretics.

United States Patent 4,438,130 discloses certain mono-oxa-, thiaspiro-cyclic-benzeneacetamide and benzamide compounds useful as analgesics.

35 One aspect of the present invention is a compound of formula



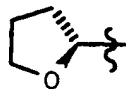
I

and the pharmaceutically acceptable acid addition salt thereof wherein R¹, R², R³, R⁴, and Ar are as defined below.

Preferred compounds of the present invention are those of formula I wherein R¹, R², R³, and R⁴ are hydrogen.

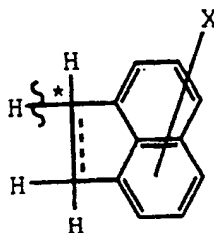
50 Other preferred compounds of the present invention are those of formula I wherein one of R¹, R², R³, and R⁴ is -OCH₃ or wherein R¹ and R⁴ are both -OCH₃.

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and R³ and R⁴ are both hydrogen.

Another aspect of the present invention is a compound of formula I wherein Ar is

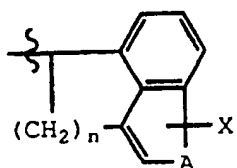


Ia

wherein X may be hydrogen, fluorine, chlorine, alkyl of from one to six carbon atoms, or aryl. Preferably X is NO₂, Cl, Br, or alkyl or alkoxy from C₁-C₆. X may be attached to the ring shown in Ia at any one of the open positions. The asterisk indicates a chiral center, both enantiomers (isomers I and II) and the racemates are preferred.

The dotted line in Ia indicates that the C₁-C₈ bond may be unsaturated or saturated.

Another aspect of the present invention is a compound of formula I wherein Ar is



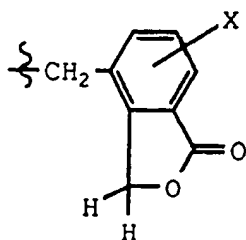
A = S or O only

n = 2 → 4 only

Ib

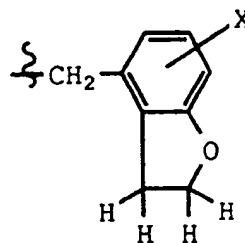
wherein n is an integer of from two to four and A is oxygen or sulfur. Preferably n is two and A is O or S. X is as above.

Another aspect of the present invention is a compound of formula I wherein Ar is



Ic

or

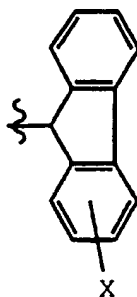


Id

wherein X is hydrogen, fluorine, chlorine, alkyl of from one to six carbon atoms, or alkoxy. Preferably X is hydrogen, Cl, Br, NO₂, or alkyl or alkoxy from C₁-C₆. X may be attached to the ring at any one of the open

positions.

Another aspect of the instant invention is a compound of formula I wherein Ar is



Ie

wherein X is hydrogen, fluorine, chlorine, alkyl of from one to six carbon atoms, or aryl. Preferably X is hydrogen, Cl, Br, or NO₂. X may be attached to the ring at any open position.

Especially preferred compounds of the present invention are:

- 1,2-dihydro-N-methyl-N-[2-(1-pyrrolidinyl)cyclohexyl]-1-acenaphthylenecarboxamide, monohydrochloride (isomer I, mixture of (1 α ,2 β) and (1 β ,2 α) forms),
- 1,2-dihydro-N-methyl-N-[2-(1-pyrrolidinyl)cyclohexyl]-1-acenaphthylenecarboxamide, monohydrochloride (isomer II, mixture of (1 α ,2 β) and (1 β ,2 α) forms),
- 25 N-[4,5-dimethoxy-2-(1-pyrrolidinyl)cyclohexyl]-1,2-dihydro-N-methyl-1-acenaphthylenecarboxamide, monohydrochloride (isomer I, mixture of (1 α ,2 β ,4 β ,5 β) and (1 β ,2 α ,4 α ,5 α) forms),
- N-[4,5-dimethoxy-2-(1-pyrrolidinyl)cyclohexyl]-1,2-dihydro-N-methyl-1-acenaphthylenecarboxamide, monohydrochloride (isomer II, mixture of (1 α ,2 β ,4 β ,5 β) and (1 β ,2 α ,4 α ,5 α) forms),
- 1,2-dihydro-N-methyl-N-[7-(1-pyrrolidinyl)-1-oxaspiro[4.5]dec-8-yl]-1-acenaphthylenecarboxamide, monohydrochloride (isomer I, (5 α ,7 α ,8 β) form), and
- 30 1,2-dihydro-N-methyl-N-[7-(1-pyrrolidinyl)-1-oxaspiro[4.5]dec-8-yl]-1-acenaphthylenecarboxamide, monohydrochloride (isomer II, (5 α ,7 α ,8 β) form).

When Isomer I and Isomer II mean the two enantiomers of the 1-acenaphthene carboxylic acid.

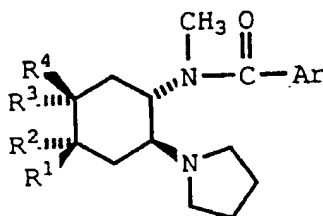
Another aspect of the present invention is a pharmaceutical composition useful as an analgesic, diuretic, antiinflammatory, neuroprotection or psychotherapeutic agent which comprises a therapeutically effective amount of a compound of formula I in combination with a pharmaceutically acceptable carrier.

Yet another aspect of the present invention is the use of a compound of formula I for the preparation of a pharmaceutical useful in alleviating pain in a mammal.

Still another aspect of the present invention is a process for the preparation of a compound of formula I described below.

DETAILED DESCRIPTION

The present invention is a novel series of 1,2-cyclohexylaminoaryl amides of formula I



I

and the pharmaceutically acceptable acid addition salts thereof wherein Ar, R¹, R², R³, and R⁴ are as described above.

The compounds of the present invention include solvates, hydrates, and pharmaceutically acceptable acid addition salts of the basic compounds of formula I above.

5 The term alkyl is a group of from one to six carbon atoms unless otherwise specified. This includes straight or branched groups such as methyl, ethyl, n-propyl, isopropyl, n-butyl, isobutyl, n-pentyl, and the like.

By virtue of the basic nitrogen on the cyclohexane moiety, pharmaceutically acceptable salts of compounds of the present invention may be prepared by reaction with appropriate acids. Suitable acids for
10 the formation of pharmaceutically acceptable salts of the compounds of this invention form a class well known to practitioners of the pharmaceutical formulation arts (of. S. M. Berge, et al., "Pharmaceutical Salts" in J. Pharm. Sci., 66: 1-19 (1977)), and include such acids as hydrochloric, hydrobromic, hydriodic, sulfuric, nitric, phosphoric, acetic, benzoic, citric, maleic, tartaric, succinic, gluconic, ascorbic, sulphamic, oxalic, pamoic, methanesulfonic, benzenesulfonic, ethanesulfonic, hydroxyethanesulfonic, and related acids and
15 mixtures thereof.

The salts are generally prepared by reacting the free base with one equivalent of the desired acid in an appropriate unreactive solvent, followed by collection of the salt by filtration or recovery upon removal of the solvent. The free base may be regenerated, if desired, by reaction of the salt with one equivalent of a base such as sodium hydroxide, sodium bicarbonate, sodium carbonate, and the like. The salts may differ from
20 the free base form of compounds of this invention in properties such as melting point and solubility in polar solvents, but are otherwise considered equivalent for the purposes of this invention.

Compounds of the present invention contain one or more asymmetric carbon atoms and exist in various stereo- and regio-isomeric forms. The present invention contemplates all stereo- and regio-isomeric forms of formula I above. The (+), (-) and (±) forms are all contemplated by the instant invention.

25 The individual stereo compounds or enantiomers are obtained, if desired, from a mixture of different forms by known methods of resolution such as the formation of diastereomers followed by recrystallization.

Compounds of the present invention are prepared by reacting an appropriate substituted cyclohexyl-diamine of formula

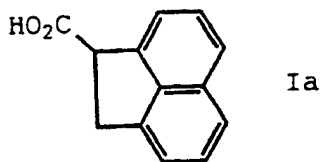


II

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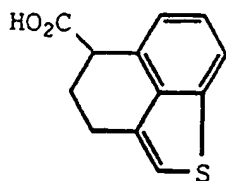
with an activated carboxylic acid

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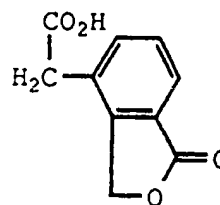


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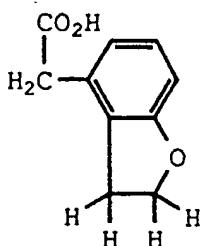
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Ib

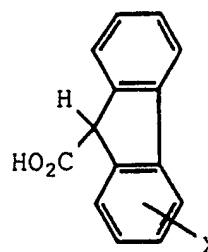


Ic



Id

or

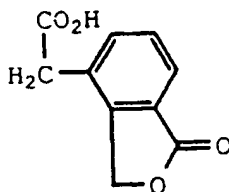


Ie

or a reactive derivative formed from such a carboxylic acid.

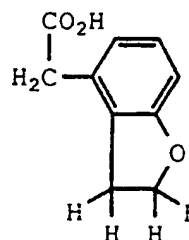
First an aryl carboxylic acid is converted to the corresponding acid halide by reaction with thionyl halide. The appropriate diamine is dissolved in a suitable solvent such as dichloromethane at low temperature. Then equimolar amounts of the acid halide and the diamine are reacted to form a compound of formula I. This is recovered by precipitation from the mixed solvent. The product can be separated into diastereomers by known means.

Compounds of the present invention wherein the aryl is



Ic

or



Id

are prepared by converting the aryl carboxylic acid into the corresponding imidazole derivative by reaction with carbonyldiimidazole. The appropriate diamine is dissolved in a suitable solvent such as dichloromethane at low temperature. Then the imidazole derivative and the diamines are reacted. The reaction mixture is quenched with water and the aqueous layer is extracted with dichloromethane.

The novel compounds of the present invention have a very high kappa opioid affinity and selectivity and potency. For example, compound 1e $K_i = 0.18$ nM with μ/κ ratio of 1416 and efficacy on rat paw pressure assay better than any known compound to the inventors with MPE_{50} 0.014 mg/kg (IV) and 0.28 mg/kg (PO).

The compounds of the present invention possess significant analgesic activity with the potential for minimum dependence liability due to their selective kappa opioid receptor properties. In addition to acting as analgesics, selective kappa opioid agonists also cause opioid receptor-mediated sedation, diuresis, and corticosteroid elevations. Accordingly, the compounds of the present invention may also be useful diuretics, antiinflammatories and psychotherapeutic agents as well as analgesics.

The compounds of the present invention also have application in congestive heart failure, advanced hepatic cirrhosis, nephrotic syndrome, chronic renal failure, trauma associated with surgery, emotional and physical stress, endocrine disorders, syndrome of inappropriate antidiuretic hormone secretion and therapy with certain pharmacologic drug agents such as certain sulfonyl ureas, clofibrate, certain tricyclics such as carbamazepine, amitriptyline, thiothixene, flubenazine and thioridazine, certain antineoplastic agents, certain analgesics, and certain natriuretic diuretics.

The compounds of the present invention also have neuroprotective indications. As such, they are useful in the treatment of stroke and the treatment of cerebral ischemia (P. F. Vonvoightlander in Brain Research, 435: 174-180 (1987)) and A. H. Tang, et al. in Brain Research, 403:52-57 (1987).

Representative examples of the compounds of this invention have shown activity in standard laboratory analgesic tests such as the rat paw pressure test as shown by the data appearing in Table 1 (M. B. Tyers, Brit. J. Pharmacol., (1980), 69, 503-512).

Moreover, representative examples of compounds of the present invention when tested *in vitro* to determine the extent of opioid receptor were found to be selectively bound to the kappa opioid receptors with much lower binding to the mu opioid receptor sites. The benefits of this selectivity in binding to opioid receptor binding sites has been discussed above and is also described in M. B. Tyers, Br. J. Pharmacol., 69:503-512 (1980).

Measurement of the kappa opioid receptor binding activity of compounds of the present invention was made by the following method. Guinea pig brain homogenates were prepared fresh daily utilizing the method of Gillan, et al., Br. J. Pharmacol., 70:481-490 (1980).

The binding of tritiated etorphine to brain homogenates was measured in the presence of unlabeled competitors compounds of the present invention with 200 nanomolar D-alanine-D-leucine-enkephalin (acronym DADLE) and 200 nanomolar D-ala-MePheGly-ol-enkephalin (acronym DAGO) added to saturate the delta and mu opioid receptors, respectively. The reaction was terminated by rapid filtration and the radioactivity bound to the filters counted by liquid scintillation spectrophotometry.

Measurement of the mu and delta opioid receptor binding activity of the compounds of this invention was made by the following method. Guinea pig brain homogenates were freshly prepared daily by the method of Gillan, et al., cited above.

Homogenates were incubated for 150 minutes at 0°C with either tritiated DAGO to measure mu receptor binding activity, or with tritiated DADLE in the presence of a tenfold excess of unlabeled DAGO to measure delta opioid receptor binding. Nonspecific binding was determined in the presence of 10⁻⁶ molar DAGO and 10⁻⁶ molar DADLE.

Reactions were terminated by rapid filtration and the radioactivity bound to the filters counted by liquid scintillation spectrophotometry.

The data were analyzed by the methods of Scatchard, Ann. N.Y. Acad. Sci., 51:660-672 (1949) and Hill, J. Physiol., 40:IV-VIII (1910). The inhibition of the binding of tritiated etorphine, DAGO and DADLE by cold ligands was determined from the regression of log percentage inhibition of specific binding or log concentration of cold ligand. The inhibition constant, K_i, was calculated from the equation:

$$K_i = \frac{IC_{50}}{1 + [L]/K_D}$$

where [L] is the concentration of the labeled ligand and K_D is the equilibrium dissociation constant.

The results of these tests are presented in Table 1.

Table 1

				Rat Paw Pressure	
	Opioid Binding k_i nM			MPE ₅₀	(mg/kg)
Compound Number	kappa	mu	mu/k	IV	PO
1a (isomer I)	16.3	2630	161	0.14	0.28
1b (isomer II)	31	387	12.5		
1c (isomer I)	4.5	578	128		
1e (isomer I)	0.18	255	1416		
1f (isomer II)	1.2	273	227		
1d (isomer II)	10.5	307	29	0.71	
3d	9.7	4930	508		
2a	0.73	653	120	0.1	
3c	5.3	2870	541	0.1	
<p>k_i values represent the mean from concentration-response curves performed in triplicate from each of at least two separate experiments.</p> <p>MPE₅₀ values represent the dose required to produce 50% of the maximum possible analgesic effect. They are derived from a single experiment with six animals per dose level.</p>					

The compounds of the present invention and/or their nontoxic, pharmaceutically acceptable acid addition salts may be administered to mammals in pharmaceutical compositions which comprise one or more compounds of this invention and/or salts thereof in combination with a pharmaceutically acceptable nontoxic carrier.

As parenteral compositions, the compounds of this invention may be administered with conventional injectable liquid carriers such as sterile, pyrogen-free water, sterile, peroxide-free ethyl oleate, dehydrated alcohols, polypropylene glycol, and mixtures thereof.

Suitable pharmaceutical adjuvants for the injectable solutions include stabilizing agents, solubilizing agents, buffers, and viscosity regulators. Examples of these adjuvants include ethanol, ethylenediamine tetraacetic acid (EDTA), tartrate buffers, citrate buffers, and high molecular weight polyethylene oxide viscosity regulators. These pharmaceutical formulations may be injected intramuscularly, intraperitoneally, or intravenously.

As solid or liquid pharmaceutical compositions, the compounds of the present invention may be administered to mammals orally in combination with conventional compatible carriers in solid or liquid form. These orally administered pharmaceutical compositions may contain conventional ingredients such as binding agents such as syrups, acacia, gelatin, sorbitol, tragacanth, polyvinylpyrrolidone, and mixtures thereof.

The compositions may further include fillers such as lactose, mannitol, starch, calcium phosphate, sorbitol, methylcellulose, and mixtures thereof.

These oral compositions may also contain lubricants such as magnesium stearate, high molecular weight polymers such as polyethylene glycol, high molecular weight fatty acids such as stearic acid, silica, or agents to facilitate disintegration of the solid formulation such as starch, and wetting agents such as sodium lauryl sulfate.

The oral pharmaceutical compositions may take any convenient form such as tablets, capsules, lozenges aqueous or oily suspensions, emulsions, or even dry powders which may be reconstituted with water or other suitable liquids prior to use.

The solid or liquid forms may contain flavorants, sweeteners, and/or preservatives such as alkyl p-hydroxybenzoates. The liquid forms may further contain suspending agents such as sorbitol, glucose, or other sugar syrups, methyl-, hydroxymethyl-, or carboxymethylcellulose, and gelatin, emulsifying agents such as lecithin or sorbitol monooleate, and conventional thickening agents. The liquid compositions may be encapsulated in, for example, gelatin capsules.

As topically administered pharmaceutical compositions, the compounds of the present invention may be

administered in the form of ointments or creams containing from about 0.1% to about 10% by weight of the active component in a pharmaceutical ointment or cream base.

Compounds of the present invention may be rectally administered in the form of suppositories. For preparing suppositories, for example, a low-melting wax such as a mixture of fatty acid glycerides or cocoa butter is first melted and the active ingredient is dispersed homogeneously in the melt. The mixture is then poured into convenient sized molds and allowed to cool and solidify.

Preferably the pharmaceutical compositions of this invention are in unit dosage form. In such form, the preparation is subdivided into unit doses containing appropriate amounts of the active component. The unit dosage can be a packaged preparation with the package containing discrete quantities of the preparation. For example, the package may take the form of packaged tablets, capsules, and powders in envelopes, vials or ampoules. The unit dosage form can also be a capsule, cachet, or tablet itself or can be the appropriate number of any of these packaged forms.

The quantity of active compound in a unit dosage form may be varied or adjusted from about 0.01 mg to about 350 mg according to the particular application and the potency of the active ingredient.

When employed systematically in therapeutic use as analgesic agents in the pharmaceutical method of this invention, the compounds are administered at doses of from about 0.0001 mg to about 2.0 mg of active compound per kilogram of body weight of the recipient.

The following examples illustrate the present invention and are not intended to be in any way limiting.

EXAMPLE 1

Preparation of 1-Acenaphthenecarboxylic Acid

1-Acenaphthenecarboxylic acid was prepared by the method of Julia [Bull. Chim. Soc. Fr., 1065 (1952)] or by the preferred, shorter route described below.

(i) 1-Tris(methylthio)methylacenaphthene

Acenaphthenol (35 g) was suspended in dry ether (200 ml) cooled to -5°C with stirring and phosphorous tribromide (8 ml) added dropwise over five minutes to give a clear solution. The solution was allowed to warm to room temperature and stirred for a further two hours. The solution was washed with water and the ether layer dried over NaCO_3 , reduced in volume at $< 40^{\circ}\text{C}$ in vacuo to 50 ml and cooled to 0°C .

Bright yellow crystals (of 1-bromoacenaphthene) were isolated by filtration and used, immediately, in the following step.

Tris(methylthio)methane (22 g) was dissolved in THF (100 ml) under N_2 at -78°C . 1.6 M BuLi in hexane solution (95 ml) was added slowly to yield a dense white precipitate. 1-Bromoacenaphthene (31.4 g) was dissolved in 20 ml of dry THF and added slowly to the above suspension to yield, after 10 minutes, a colorless solution. This was allowed to warm to room temperature, with the development of a brown color, and stirred overnight.

This solution was evaporated in vacuo to yield a brown oil (48 g). This material was crystallized from hexane to give orange crystals (29 g) of 1-(trimethylthio)methyl acenaphthene. m.p. $68-70^{\circ}\text{C}$

Found: C, 62.62; H, 5.87; S, 31.11;

Theory: C, 62.70; H, 5.92; S, 31.62;

(ii) Methyl 1-acenaphthenecarboxylate

1-Tris(methylthio)methyl acenaphthene (18.4 g), Mercury (II) Chloride (68 g) and Mercury (II) Oxide (22 g) were slurried in $\text{MeOH}/\text{H}_2\text{O}$, 12/1 (1500 ml), stirred at room temperature for 18 hours, then refluxed for two hours, and then allowed to cool.

The suspension was filtered and the cake washed with dichloromethane (2 x 100 ml). Water (1500 ml) was added to the solution and the total extracted with dichloromethane (2 x 1 L). The dichloromethane

solution was washed with 75% aqueous ammonium acetate (2 x 500 ml), dried over MgSO_4 and evaporated in vacuo to yield an orange oil (14 g). This was then purified by chromatography using hexane/ether 25/1 as eluent and Merck 15/11 silica gel as stationary phase to give a yellow liquid (10 g).

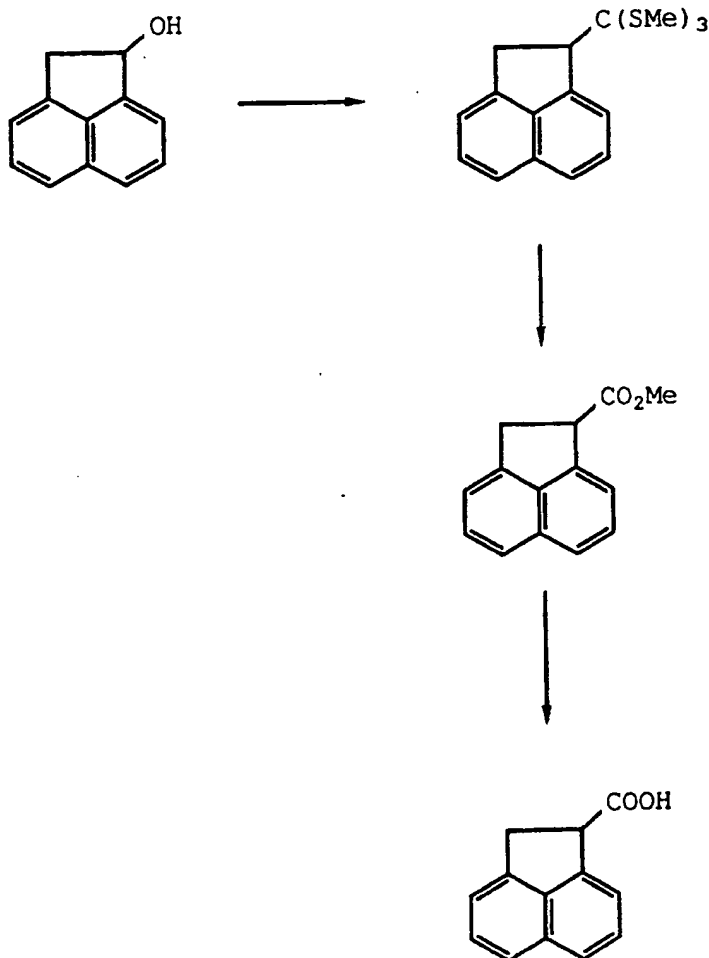
NMR 300 MHz, CDCl_3 : δ 3.67 (dd) 1H, 3.78 (s) 3H, 3.91 (dd) 1H, 4.62 (dd) 1H, 7.35 (m) 1H, 7.5 (m) 3H, 7.7 (m) 2H.

(iii) 1-Acenaphthenecarboxylic acid

The above ester (1.9 g) was dissolved in 5 ml THF and this solution added to 6N KOH (40 ml). The mixture was refluxed for 10 hours and allowed to cool, then acidified with 5N HCl solution and extracted with ethyl acetate (3 x 50 ml). The extracts were combined, dried over MgSO_4 and evaporated in vacuo to yield a gummy orange solid (1.6 g). This was recrystallized from hexane/toluene 3/1 to yield orange crystals, mp 138-145°C.

Scheme

Alternative Preferred Route to Acenaphthene
Carboxylic Acid



The acenaphthenecarboxylic acid (1.1 mmol) was stirred in neat thionyl chloride (5 ml) at reflux for one hour. The thionyl chloride was evaporated in vacuo and the resulting oil treated with 25 ml CCl_4 . This

mixture was then evaporated in vacuo to yield a brown oil which was used without further purification.

The appropriate diamine of structure II (1 mmol) was dissolved in a small volume (5 ml) of dichloromethane and cooled to 0 °C with stirring. The brown oil (above) was also dissolved in a similar volume of dichloromethane at room temperature and added to the acid solution dropwise. Stirring was continued for one hour at room temperature. The solution was then triturated with Et₂O to yield a precipitate which was isolated by filtration.

This product was then separated into diastereoisomers (racemic pairs for Ia, Ib, Ic, Id and enantiomers for Ie and If using medium pressure silica gel chromatography (Merck 11695 15 µm silica, CH₂Cl₂/MeOH 10:1 eluant). The appropriate fractions were collected, evaporated in vacuo, dissolved in dichloromethane (5 ml). This solution was then treated with ethereal HCl and evaporated in vacuo to yield a white solid, recrystallized from isopropanol.

1.a. 1,2-dihydro-N-methyl-N-[2-(1-pyrrolidinyl)cyclohexyl]-1-acenaphthylenecarboxamide, monohydrochloride (isomer I, mixture of (1 α ,2 β) and (1 β , 2 α) forms)

Elemental analysis for C₂₄H₃₀N₂OHCl·0.4H₂O requires:

Calcd: C, 71.00; H, 7.89; N, 6.90

Found: C, 71.01; H, 7.75; N, 6.92

NMR 300 MHz (DMSO-d₆) δ 1 -> 2 (m) 12H, 3.20 (s) 3H, 3.30 (m) 4H, 3.48 (m) 1H, 3.65 (m) 1H, 3.78 (d of d) 1H, 4.55 (m) 1H, 5.02 (d of d) 1H, 7.4 -> 7.6 (m) 5H.

mp > 230 °C.

IR (liquid film) 1641 cm⁻¹ NHCO; 3400 cm⁻¹ \dot{N} H

1.b. 1,2-dihydro-N-methyl-N-[2-(1-pyrrolidinyl)cyclohexyl]-1-acenaphthylenecarboxamide, monohydrochloride (isomer II, mixture of (1 α ,2 β) and (1 β and 2 α) forms)

Elemental analysis for C₂₄H₃₀N₂OHCl·0.15H₂O requires:

Calcd: C, 71.80; H, 7.85; N, 6.98

Found: C, 71.79; H, 7.84; N, 7.02

NMR 300 MHz (DMSO-d₆) δ 1 -> 2 (m) 12H, 3.20 (m) and 3.32 (s) integral obscured, 3.69 (m) 4H, 4.55 (m) 1H, 5.02 (d of d) 1H, 7.22 (d) 1H, 7.32, 1H, 7.6 (m) 4H.

mp 228-232 °C.

IR (liquid film) 1641 cm⁻¹ NHCO; 3413 \dot{N} -H

1.c. N-[4,5-dimethoxy-2-(1-pyrrolidinyl)cyclohexyl]-1,2-dihydro-N-methyl-1-acenaphthylenecarboxamide, monohydrochloride (isomer I, mixture of (1 α ,2 β ,4 β ,5 β) and (1 β ,2 α ,4 α ,5 α) forms)

Elemental analysis for C₂₆H₃₄N₂O₃·HCl requires:

Calcd: C, 68.03; H, 7.69; N, 6.10

Found: C, 67.88; H, 7.72; N, 6.07

NMR 300 MHz (DMSO-d₆) δ 1.5 -> 2.2 (m) 8H, 3.1 (m) 1H, 3.18 (s) 3H, 3.30 (m) and 3.30 (s) and 3.35 (s) Integral obscured, 3.50 (m) 1H, 3.75 (m) 3H, 4.30 (m) 1H, 5.02 (d of d) 1H, 7.22 (d) 1H, 7.55 (m) 5H.

mp 137-140 °C.

IR liquid film 1637 cm⁻¹; NHCO 3402 cm⁻¹ \dot{N} -H

1.d. N-[4,5-dimethoxy-2-(1-pyrrolidinyl)cyclohexyl]-1,2-dihydro-N-methyl-1-acenaphthylenecarboxamide, monohydrochloride (isomer II, mixture of (1 α ,2 β , 4 β ,5 β) and (1 β ,2 α ,4 α ,5 α) forms)

Elemental analysis for C₂₆H₃₄N₂O₃HCl·H₂O requires:

Calcd: C, 65.46; H, 7.82; N, 5.87

Found: C, 65.47; H, 7.82; N, 5.90

NMR 300 MHz (DMSO-d₆) δ 1.5 -> 2.2 (m) 8H, 3.1 (m) 1H, 3.28 (s) and 3.30 (s) and 3.33 (s) and 3.50 (m) Integral obscured, 3.80 (m) 3H, 4.80 (m) 1H, 5.02 (d of d) 1H, 7.22 (d) 1H, 7.31 (d) 1H, 7.61 (m) 4H.

mp 126-133 °C.

IR Liquid film 1636 cm⁻¹ NHCO; 3392 cm⁻¹ \dot{N} -H

1.e. 1,2-dihydro-N-methyl-N-[7-(1-pyrrolidinyl)-1-oxaspiro[4.5]dec-8-yl]-1-acenaphthylenecarboxamide, monohydrochloride (isomer I, (5 α ,7 α ,8 β) form)

Elemental analysis for C₂₇H₃₄N₂O₂HCl·0.5H₂O requires:

5 Calcd: C, 69.88; H, 7.81; N, 6.03

Found: C, 69.84; H, 7.77; N, 5.94

NMR 300 MHz (DMSO-d₆) δ 1.5 -> 2 (m) 14H, 3.18 (s) 3H, 3.25 (m) Integral obscured 3.52 (m) 1H, 3.75 (m) 5H, 4.52 (m) 1H, 5.02 (d of d) 1H, 7.25 (d) 1H, 7.55 (m) 5H.

mp 143-146 °C.

10 $[\alpha]_D^{20} = +13.9^\circ$, C = 0.17, CH₂Cl₂

IR Liquid film 1640 cm⁻¹; NHCO 3392 cm⁻¹ \hat{N} H

1.f. 1,2-dihydro-N-methyl-N-[7-(1-pyrrolidinyl)-1-oxaspiro[4.5]dec-8-yl]-1-acenaphthylenecarboxamide, monohydrochloride (isomer II, (5 α ,7 α ,8 β) form)

15 Elemental analysis for C₂₇H₃₄N₂O₂HCl·1.3H₂O requires:

Calcd: C, 67.77; H, 7.92; N, 5.84

Found: C, 67.50; H, 8.17; N, 5.48

NMR 300 MHz (DMSO-d₆) δ 1.5 -> 2.1 (m) 14H, 3.20 (s) and 3.4 -> 3.8 (m) Integral obscured, 4.55 (m) 1H, 5.02 (d of d) 1H, 7.21 (d) 1H, 7.32 (d) 1H, 7.55 (m) 4H.

mp 138-140 °C.

20 $[\alpha]_D^{20} = +43.5^\circ$ C = 0.54, CH₂Cl₂

IR Liquid film 1641 cm⁻¹ NHCO; 3398 cm⁻¹ \hat{N} -H

25 EXAMPLE 2

The 9-fluorenyl carboxylic acid (1.1 mmol) was stirred in neat thionyl chloride at reflux for one hour. The thionyl chloride was evaporated in *vacuo* and the resulting oil treated with carbon tetrachloride (25 ml). This mixture was then evaporated in *vacuo* to yield a brown oil (not isolated).

30 The spiroether diamine (1 mmol) was dissolved in a small volume (5 ml) of dichloromethane at 0 °C with stirring. The acid chloride from above was dissolved in a similar volume of dichloromethane and this solution added to the amine solution dropwise.

Stirring was continued for one hour at room temperature, after which time the solution was triturated with H₂O to yield an off-white precipitate. Recrystallized from propan-2-ol to yield white solid.

2.a. (-)-(5 α ,7 α ,8 β)-N-methyl-N-[7-(1-pyrrolidinyl)-1-oxaspiro[4.5]dec-8-yl]-9H-fluorene-9-carboxamide, monohydrochloride

40 Elemental analysis for C₂₈H₃₄N₂O₂HCl·0.7H₂O requires:

Calcd: C, 70.06; H, 7.51; N, 5.79

Found: C, 70.06; H, 7.65; N, 5.84

NMR 300 MHz (DMSO-d₆) δ 1.5 -> 2.1 (m) 14H, 3.1 -> 3.8 (m) + 3.32 (s) Integral obscured, 4.55 (broad m) 1H, 5.49 (s) 1H, 7.35 (m) and 7.90 (m) 8H.

mp 201-203 °C.

45 IR (liquid film) 1641 cm⁻¹ NHCO 3401 \hat{N} -H

2.b. (±)-trans-N-methyl-N-[2-(1-pyrrolidinyl)cyclohexyl]-9H-fluorene-9-carboxamide, monohydrochloride

50 Elemental analysis for C₂₅H₃₀N₂O·HCl·1.4H₂O requires:

Calcd: C, 68.73; H, 7.81; N, 6.41

Found: C, 68.73; H, 7.93; N, 6.25

mp 243-252 ° (iPrOH) (white microcrystalline solid).

IR (neat) 3416, 1641 cm⁻¹

Mass Spec: (EI) m/e 374 (M⁺, 5%) 151 (100%)

NMR δ (DMSO-d₆) 300 MHz 10.24 (1H, br, s), 7.95 (3H, m), 7.35 (5H, m), 5.45 (1H, s), 4.58 (1H, m), 3.72 (1H, m), 3.47 (3H, s), 3.35 (4H, m), 2.2-1.22 (12H).

EXAMPLE 3

4-Isobenzofuran-1-one acetic acid (1 mmol) and carbonyldiimidazole (1.1 mmol) were stirred under
 5 reflux in THF (10 ml) for one hour. The THF was then evaporated in vacuo and the residual oil dissolved in
 a small volume of dichloromethane (25 ml).

The diamine was dissolved in a similar small volume of dichloromethane (25 ml), cooled to 0 °C and the
 above solution added dropwise to it, with stirring. The resulting mixture was stirred at room temperature for
 a further two hours and then quenched with water. The aqueous layer was extracted with dichloromethane
 10 (3 x 100 ml), the extracts combined, washed with water, dried over MgSO₄, and evaporated in vacuo.

The resulting oil was chromatographed using Merck Silica Gel Art 11695, CH₂Cl₂/MeOH 10:1 on a
 medium pressure preparative system.

The appropriate fractions were evaporated, the residual oil dissolved in 5 ml of dichloromethane and
 treated with ethereal HCl to yield a crystalline white material which was recrystallized from propan-2-ol.

15

3a. (±)-trans-1,3-dihydro-N-methyl-1-oxo-N-[2-(1-pyrrolidinyl)cyclohexyl]-4-isobenzofuranacetamide, mon-
hydrochloride

20 Elemental analysis for C₂₁H₂₈N₂O₃ · HCl 0.3H₂O requires:

Calcd: C, 63.32; H, 7.49; N, 7.03

Found: C, 63.33; H, 7.37; N, 7.01

NMR 300 MHz (DMSO-d₆) δ 1.2 -> 2.0 (m) 12H, 3.05 (s) 3H, 3.55 (m) 2H, 3.74 (d) 1H, 4.20 (d) 1H, 4.55
 (broad m) 1H, 5.37 (d) 1H, 5.42 (d) 1H, 7.53 (t) 1H, 7.63 (d) 1H, 7.72 (d) 1H.

25 mp 251-254 °C.

IR (liquid film) 1645 cm⁻¹ NHCO 1745 cm⁻¹ OCO, 3362 cm⁻¹ $\dot{N}H$

3b. (±)-(1 α ,2 β ,4 β ,5 β)-N-[4,5-dimethoxy-2-(1-pyrrolidinyl) cyclohexyl]-1,3-dihydro-N-methyl-1-oxo-4-isoben-
zofuranacetamide mixture of (1 α ,2 β ,4 β ,5 β) and (1 β ,2 α ,4 α ,5 α) forms)

30

Elemental analysis for C₂₃H₃₂N₂O₅ · HCl requires:

Calcd: C, 60.99; H, 7.34; N, 6.18

Found: C, 60.75; H, 7.41; N, 6.20

NMR 300 MHz (DMSO-d₆) δ 1.6 -> 2.0 (m) 8H, 3.05 (s) and 3.10 (m) 4H, 3.35 (m) and 3.31 (s) and 3.33 (s)
 35 3.35 (m) Integral obscured, 3.52 (m) 1H, 3.75 (m) 2H, 3.82 (d) 1H, 4.24 (d) 1H, 4.80 (broad m) 1H, 5.38 (d)
 1H, 7.52 (t) 1H, 7.62 (d) 1H, 7.71 (d) 1H.

mp 250-252 °C. IR (liquid film) 1646 cm⁻¹ NHCO; 1757 cm⁻¹ COO; 3419 cm⁻¹ $\dot{N}H$.

3c. (-)-(5 α ,7 α ,8 β)-1,3-dihydro-N-methyl-1-oxo-N-[7-(1-pyrrolidinyl)-1-oxaspiro[4.5]dec-8-yl]-4-isoben-
 40 zofuranacetamide monohydrochloride

Elemental analysis for C₂₄H₃₂N₂O₄ · HCl · H₂O requires:

Calcd: C, 61.73; H, 7.55; N, 6.00; Cl, 7.59

Found: C, 61.99; H, 7.52; N, 5.91; Cl, 7.38

NMR 300 MHz (DMSO-d₆) δ 1.5 -> 2.0 (m) 14H, 3.05 (s) 3H, 3.2 -> 3.6 (m) Integral obscured, 3.86 (d) 1H,
 45 4.19 (d) 1H, 4.58 (broad m) 1H, 5.35 (d) 1H, 5.43 (d) 1H, 7.55 (t) 1H, 7.66 (d) 1H, 7.75 (d) 1H.

mp 182-184 °C.

IR (liquid film) 1645 cm⁻¹ NHCO; 1757 cm⁻¹ OCO; 3425 cm⁻¹ $\dot{N}H$

$[\alpha]_D^{20}$ = -1.71 °C = 0.5 CH₂Cl₂

50

3d. (-)-(5 α ,7 α ,8 β)-2,3-dihydro-N-methyl-N-[7-(1-pyrrolidinyl)-1-oxaspiro[4.5]dec-8-yl]-4-benzofuranacetamide
monohydrochloride

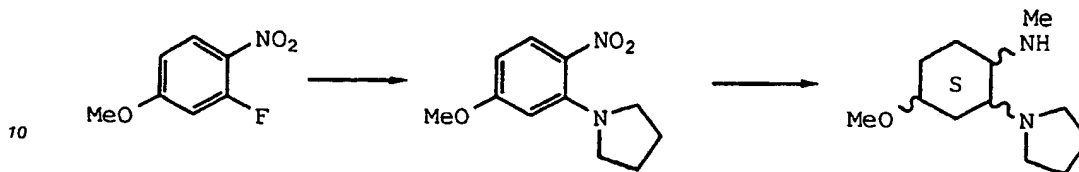
IR max: 1646 (s), 1597 cm⁻¹

55 $[\alpha]_D$ = -20 °, ϵ = 0.64 (CH₂Cl₂).

EXAMPLE 4

A method for preparing the diamine intermediates to 1,2-dihydro-N-[4-methoxy-2-(1-pyrrolidinyl)-cyclohexyl]-N-methyl-1-acenaphthylenecarboxamide monohydrochloride (isomers I and II, mixtures of (1 α ,2 β ,4 α) forms), and 1,2-dihydro-N-[4-methoxy-2-(1-pyrrolidinyl)cyclohexyl]-N-methyl-1-acenaphthylenecarboxamide, monohydrochloride (isomers I and II, mixtures of (1 α ,2 β ,4 β) forms).

5



10

15

3-Fluoro-4-nitroanisole

3-Fluoro-4-nitrophenol (Fluorochem) (10 g, 64 mmol) was dissolved in butan-2-one (60 ml) and treated with potassium carbonate 16.6 g (120 mmol) at 40 °C for 10 minutes. The resulting suspension was cooled to 0 °C, treated with methyl iodide (7.5 ml, 120 mmol), heated to 40 °C for three hours and concentrated in vacuo to 20 ml. The mixture was poured into dichloromethane (35 ml), filtered and the filtrate evaporated to give 3-fluoro-4-nitroanisole, as a white solid (8.0 g, 73%). An analytically pure sample was obtained by recrystallization from aqueous ethanol (4:1); mp 47-49 °C; ir 1608 cm⁻¹.

20

25

3-(1-Pyrrolidinyl)-4-nitroanisole

3-Fluoro-4-nitroanisole (8.0 g, 47 mmol) was added over 10 minutes to pyrrolidine (25 ml) at room temperature. The resulting mixture was poured into dilute aqueous sodium hydroxide (100 ml) and extracted with dichloromethane (4 x 50 ml) to give an orange solid (7.5 g) which was purified by silica gel chromatography (CH₂Cl₂ eluant) to give 3-(1-pyrrolidinyl)-4-nitroanisole as an orange solid (5.2 g, 50%), mp 46-48.5 °C; ir 1613, 1569 cm⁻¹.

30

35

Hydrogenation of 3-pyrrolidinyl-4-nitroanisole

3-(1-Pyrrolidinyl)-4-nitroanisole (0.50 g, 2.2 mmol), 5% rhodium on alumina (0.40 g), isopropyl alcohol (75 ml) and 48 wt percent aqueous fluoroboric acid (80 mg) were treated with hydrogen at 1000 psi and 80 °C for eight hours. The resulting mixture was filtered, concentrated in vacuo, dissolved in ethyl formate (5 ml) and triethylamine (1 ml) and heated to reflux for 50 minutes. The mixture was concentrated in vacuo and the residue dissolved in tetrahydrofuran (5 ml) then treated with a 1.0 M solution of lithium aluminum hydride in diethyl ether (4 ml) at 30-40 °C for 12 hours. Aqueous sodium hydroxide (0.2 ml) was added and the resulting precipitate removed by filtration. The filtrate was evaporated to give an oil (0.9 g) which contains (±)-(1 α ,2 α ,4 α)-4-methoxy-N-methyl-2-(1-pyrrolidinyl)cyclohexanamine, (±)-(1 α ,2 α ,4 β)-4-methoxy-N-methyl-2-(1-pyrrolidinyl)cyclohexanamine, (±)-(1 α ,2 β ,4 α)-4-methoxy-N-methyl-2-(1-pyrrolidinyl)cyclohexanamine, and (±)-(1 α ,2 β ,4 β)-4-methoxy-N-methyl-2-(1-pyrrolidinyl)cyclohexanamine.

40

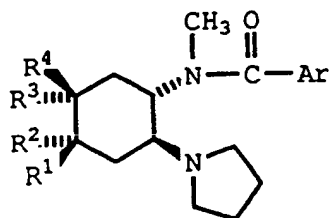
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Claims

50

1. A compound of formula

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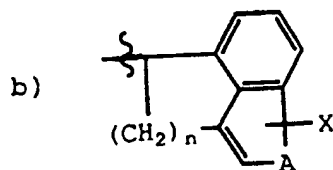
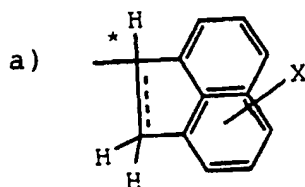


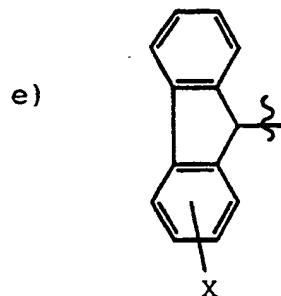
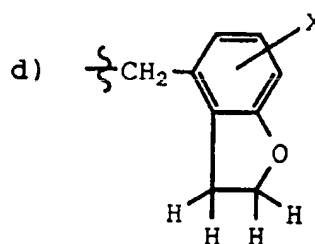
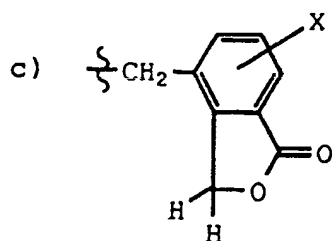
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- and the pharmaceutically acceptable acid addition salts thereof
 wherein $R_1, R_2, R_3, R_4 = H$; or
 wherein one of R_1, R_2, R_3, R_4 may be $= OCH_3$ when the rest are all H ; or
 wherein $R_1 = R_4 = OCH_3$, and $R_2 = R_3 = H$;
 R^1 and R^2 together form



- and R_3, R_4 are H ;
 Ar is





wherein X is hydrogen, fluorine, chlorine, alkyl of from one to six carbon atoms or aryl,
n is an integer from 2 to 4, and

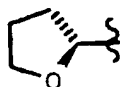
A is oxygen or sulfur.

2. A compound according to Claim 1 wherein R¹, R², R³, and R⁴ are hydrogen.

3. A compound according to Claim 1 where one of R¹, R², R³, or R⁴ is OCH₃.

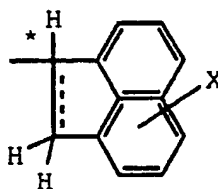
4. A compound according to Claim 1 where R¹ and R⁴ are both -OCH₃.

5. A compound according to Claim 1 wherein R¹ and R² taken together form



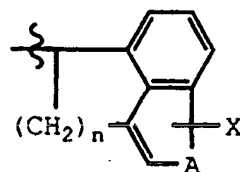
and R³ and R⁴ are hydrogen.

6. A compound according to Claim 1 wherein Ar is



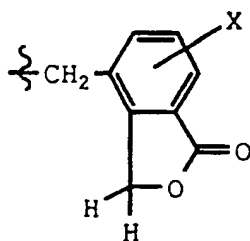
(a)

7. A compound according to Claim 1 wherein Ar is



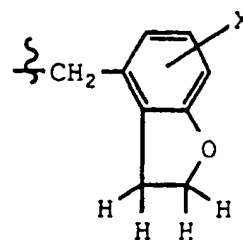
(b)

8. A compound according to Claim 1 wherein Ar is



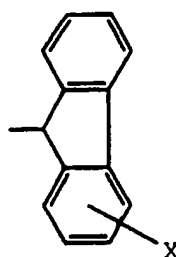
(1c)

or



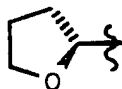
(1d)

9. A compound according to Claim 1 wherein Ar is



(e)

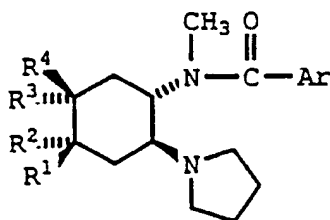
and R_1 , R_2 , R_3 , and R_4 are hydrogen or R_3 and R_4 are hydrogen and R_1 and R_2 form



and R^3 and R^4 are hydrogen.

10. A compound according to Claim 1 selected from the group consisting of:
- 1,2-dihydro-N-methyl-N-[2-(1-pyrrolidinyl)cyclohexyl]-1-acenaphthylenecarboxamide, monohydrochloride (isomer I, mixture of $(1\alpha,2\beta)$ and $(1\beta,2\alpha)$ forms),
 - 1,2-dihydro-N-methyl-N-[2-(1-pyrrolidinyl)cyclohexyl]-1-acenaphthylenecarboxamide, monohydrochloride (isomer II, mixture of $(1\alpha,2\beta)$ and $(1\beta,2\alpha)$ forms),
 - N-[4,5-dimethoxy-2-(1-pyrrolidinyl)cyclohexyl]-1,2-dihydro-N-methyl-1-acenaphthylenecarboxamide, monohydrochloride (isomer I, mixture of $(1\alpha,2\beta,4\beta,5\beta)$ and $(1\beta,2\alpha,4\alpha,5\alpha)$ forms),
 - N-[4,5-dimethoxy-2-(1-pyrrolidinyl)cyclohexyl]-1,2-dihydro-N-methyl-1-acenaphthylenecarboxamide, monohydrochloride (isomer II, mixture of $(1\alpha,2\beta,4\beta,5\beta)$ and $(1\beta,2\alpha,4\alpha,5\alpha)$ forms),

- 1,2-dihydro-N-methyl-N-[7-(1-pyrrolidinyl)-1-oxaspiro[4.5]dec-8-yl]-1-acenaphthylenecarboxamide, monohydrochloride (isomer I, (5 α ,7 α ,8 β) form),
 1,2-dihydro-N-methyl-N-[7-(1-pyrrolidinyl)-1-oxaspiro[4.5]dec-8-yl]-1-acenaphthylenecarboxamide, monohydrochloride (isomer II, (5 α ,7 α ,8 β) form),
 5 1,2-dihydro-N-[4-methoxy-2-(1-pyrrolidinyl)cyclohexyl]-N-methyl-1-acenaphthylenecarboxamide, monohydrochloride (isomers I and II, mixtures of (1 α ,2 β ,4 β) and (1 β ,2 α ,4 α) forms),
 1,2-dihydro-N-[4-methoxy-2-(1-pyrrolidinyl)cyclohexyl]-N-methyl-1-acenaphthylenecarboxamide, monohydrochloride (isomers I and II, mixtures of (1 β ,2 α ,4 α) and (1 α ,2 β ,4 β) forms).
 11. A compound according to Claim 1 selected from the group consisting of:
 10 (-)-(5 α ,7 α ,8 β)-N-methyl-N-[7-(1-pyrrolidinyl)-1-oxaspiro[4.5]dec-8-yl]-9H-fluorene-9-carboxamide, monohydrochloride and,
 (±)-N-methyl-N-[2-(1-pyrrolidinyl)cyclohexyl]-9H-fluorene-9-carboxamide, monohydrochloride.
 12. A compound according to Claim 1 selected from the group consisting of:
 (±)-trans-1,3-dihydro-N-methyl-1-oxo-N-[2-(1-pyrrolidinyl)cyclohexyl]-4-isobenzofuranacetamide, monohydrochloride,
 15 (±)-(1 α ,2 β ,4 β ,5 β)-N-[4,5-dimethoxy-2-(1-pyrrolidinyl)cyclohexyl]-1,3-dihydro-N-methyl-1-oxo-4-isobenzofuranacetamide,
 (-)-(5 α ,7 α ,8 β)-1,3-dihydro-N-methyl-1-oxo-N-[7-(1-pyrrolidinyl)-1-oxaspiro[4.5]dec-8-yl]-4-isobenzofuranacetamide monohydrochloride, (-)-(5 α ,7 α ,8 β)-2,3-dihydro-N-methyl-N-[7-(1-pyrrolidinyl)-1-oxaspiro[4.5]dec-8-yl]-4-benzofuranacetamide monohydrochloride.
 20 13. A pharmaceutical composition comprising a therapeutically effective amount of a compound according to Claim 1 to 12 in combination with a pharmaceutically acceptable carrier.
 14. Use of a compound of formula I according to claims 1 to 12 for the preparation of a pharmaceutical composition useful for alleviating pain, increasing urination, treating inflammation, or treating psychotherapeutic disorders in a mammal.
 25 15. A process for the preparation of a compound of formula I



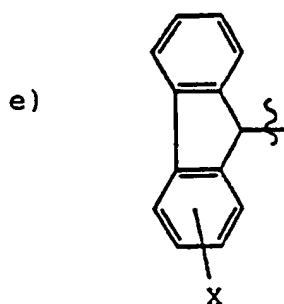
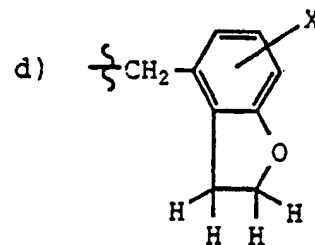
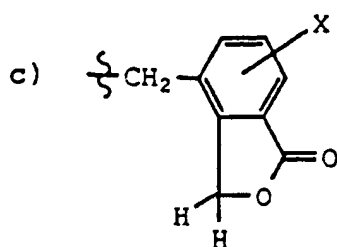
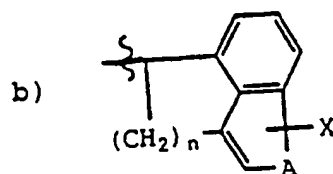
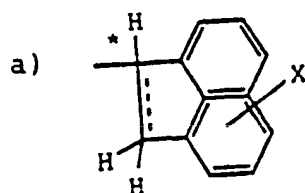
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- wherein R₁, R₂, R₃, R₄ = H; or
 40 wherein one of R₁, R₂, R₃, R₄ may be = OCH₃ when the rest are all H; or
 wherein R₁ = R₄ = OCH₃, and R₂ = R₃ = H;
 R₁ and R₂ together form



- 50 and R₃, R₄ are H;
 Ar is

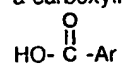
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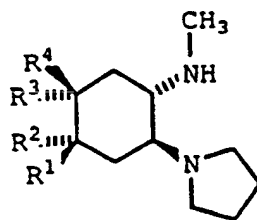
wherein X is hydrogen, fluorine, chlorine, alkyl of from one to six carbon atoms or aryl,
n is an integer from 2 to 4, and

A is oxygen or sulfur
characterized in that if

I. Ar is a) b.) or e.)
a carboxylic acid of formula



is converted to the corresponding acid chloride by reaction with thionyl chloride, which is reacted with a
diamine of the formula II

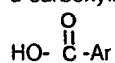


II

wherein R¹ R² R³ and R⁴ have the above meaning and if

II. Ar is c.) or d.)

a carboxylic acid of formula

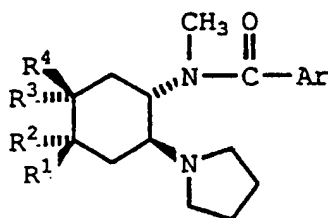


is converted to the corresponding imidazole by reaction with carbonyldiimidazole, which is reacted with a diamine of formula II above

whereafter the compound of formula I if desired is converted to a pharmaceutically acceptable salt thereof.

Claims for the following Contracting State: ES

1. A process for the preparation of a compound of formula I



I

and the pharmaceutically acceptable acid addition salts thereof

wherein R₁, R₂, R₃, R₄ = H; or

wherein one of R₁, R₂, R₃, R₄ may be = OCH₃ when the rest are all H; or

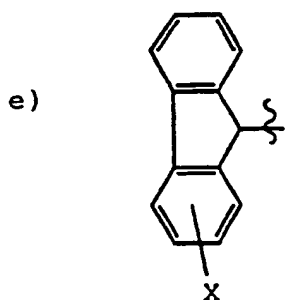
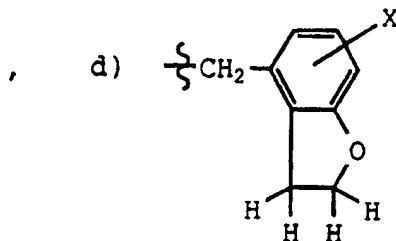
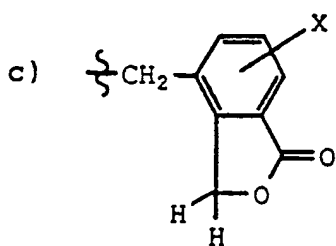
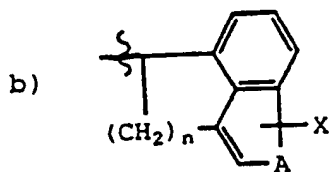
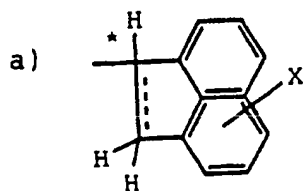
wherein R₁ = R₄ = OCH₃, and R₂ = R₃ = H;

R₁ and R₂ together form



and R₃, R₄ are H;

Ar is



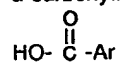
wherein X is hydrogen, fluorine, chlorine, alkyl of from one to six carbon atoms or aryl,

n is an integer from 2 to 4, and

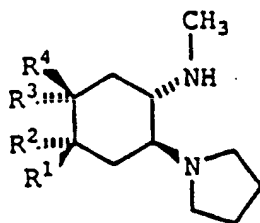
A is oxygen or sulfur
characterized in that if

I. Ar is a) b.) or e.)

a carboxylic acid of formula



is converted to the corresponding acid chloride by reaction with thionyl chloride, which is reacted with a
diamine of the formula II

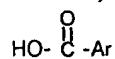


II

wherein R¹ R² R³ and R⁴ have the above meaning and if

II. Ar is c.) or d.)

a carboxylic acid of formula



is converted to the corresponding imidazol by reaction with carbonyldiimidazole, which is reacted with a diamine of formula II above

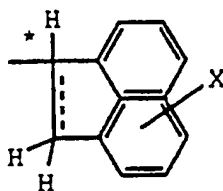
whereafter the compound of formula I if desired is converted to a pharmaceutically acceptable salt thereof.

2. A process for preparing a compound according to Claim 1 wherein R¹ R², R³, and R⁴ are hydrogen.
3. A process for preparing a compound according to Claim 1 where one of R¹ R², R³, or R⁴ is OCH₃.
4. A process for preparing a compound according to Claim 1 where R¹ and R⁴ are both -OCH₃.
5. A process for preparing a compound according to Claim 1 wherein R¹ and R² taken together form



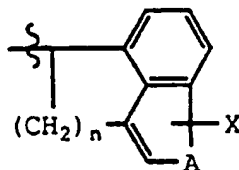
and R³ and R⁴ are hydrogen.

6. A process for preparing a compound according to Claim 1 wherein Ar is



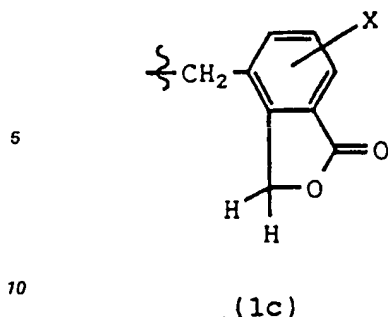
(a)

7. A process for preparing a compound according to Claim 1 wherein Ar is

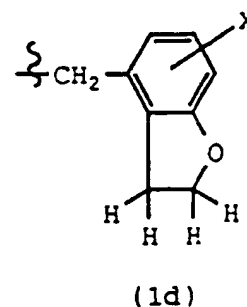


(b)

8. A process for preparing a compound according to Claim 1 wherein Ar is



or



15 9. A process for preparing a compound according to Claim 1 wherein Ar is



and R₁, R₂, R₃, and R₄ are hydrogen or R₃ and R₄ are hydrogen and R₁ and R₂ form



35 and R³ and R⁴ are hydrogen.

10. A process for preparing a compound according to Claim 1 selected from the group consisting of:
- 1,2-dihydro-N-methyl-N-[2-(1-pyrrolidinyl)cyclohexyl]-1-acenaphthylenecarboxamide, monohydrochloride (isomer I, mixture of (1 α ,2 β) and (1 β ,2 α) forms),
 - 1,2-dihydro-N-methyl-N-[2-(1-pyrrolidinyl)cyclohexyl]-1-acenaphthylenecarboxamide, monohydrochloride (isomer II, mixture of (1 α ,2 β) and (1 β ,2 α) forms),
 - N-[4,5-dimethoxy-2-(1-pyrrolidinyl)cyclohexyl]-1,2-dihydro-N-methyl-1-acenaphthylenecarboxamide, monohydrochloride (isomer I, mixture of (1 α ,2 β ,4 β ,5 β) and (1 β ,2 α ,4 α ,5 α) forms),
 - N-[4,5-dimethoxy-2-(1-pyrrolidinyl)cyclohexyl]-1,2-dihydro-N-methyl-1-acenaphthylenecarboxamide, monohydrochloride (isomer II, mixture of (1 α ,2 β ,4 β ,5 β) and (1 β ,2 α ,4 α ,5 α) forms),
 - 1,2-dihydro-N-methyl-N-[7-(1-pyrrolidinyl)-1-oxaspiro[4.5]dec-8-yl]-1-acenaphthylenecarboxamide, monohydrochloride (isomer I, (5 α ,7 α ,7 β) form),
 - 1,2-dihydro-N-methyl-N-[7-(1-pyrrolidinyl)-1-oxaspiro[4.5]dec-8-yl]-1-acenaphthylenecarboxamide, monohydrochloride (isomer II, (5 α ,7 α ,8 β) form),
 - 1,2-dihydro-N-[4-methoxy-2-(1-pyrrolidinyl)cyclohexyl]-N-methyl-1-acenaphthylenecarboxamide, monohydrochloride (isomers I and II, mixtures of (1 α ,2 β ,4 β) and (1 β ,2 α ,4 α) forms),
 - 1,2-dihydro-N-[4-methoxy-2-(1-pyrrolidinyl)cyclohexyl]-N-methyl-1-acenaphthylenecarboxamide, monohydrochloride (isomers I and II, mixtures of (1 β ,2 α ,4 α) and (1 α ,2 β ,4 β) forms).
11. A process for preparing a compound according to Claim 1 selected from the group consisting of:
- (-)-(5 α ,7 α ,8 β)-N-methyl-N-[7-(1-pyrrolidinyl)-1-oxaspiro[4.5]dec-8-yl]-9H-fluorene-9-carboxamide, monohydrochloride and,
 - (\pm)-trans-N-methyl-N-[2-(1-pyrrolidinyl)cyclohexyl]-9H-fluorene-9-carboxamide, monohydrochloride.
12. A process for preparing a compound according to Claim 1 selected from the group consisting of:
- (\pm)-trans-1,3-dihydro-N-methyl-1-oxo-N-[2-(1-pyrrolidinyl)cyclohexyl]-4-isobenzofuranacetamide, monohydrochloride

ohydrochloride,

(±)-(1 α ,2 β ,4 β ,5 β)-N-[4,5-dimethoxy-2-(1-pyrrolidinyl)cyclohexyl]1,3-dihydro-N-methyl-1-oxo-4-isobenzofuranacetamide,

(-)-(5 α ,7 α ,8 β)-1,3-dihydro-N-methyl-1-oxo-N-[7-(1-pyrrolidinyl)-1-oxaspiro[4.5]dec-8-yl]-4-

5 isobenzofuranacetamide monohydrochloride, (-)-(5 α ,7 α ,8 β)-2,3-dihydro-N-methyl-N-[7-(1-pyrrolidinyl)-1-oxaspiro[4.5]dec-8-yl]-4-benzofuranacetamide monohydrochloride.

13. Use of a compound of formula I prepared according to claims 1 to 12 for the preparation of a pharmaceutical composition useful for alleviating pain, increasing urination, treating inflammation, or treating psychotherapeutic disorders in a mammal.

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European Patent
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EUROPEAN SEARCH REPORT

Application Number

EP 90 10 1339

DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int. Cl.5)
A	EP-A-0 147 085 (WARNER-LAMBERT CO.) * pages 3,4 *; & US - A - 4 656 182 (Cat. D) ---	1,13	C 07 D 295/135 C 07 D 405/08 A 61 K 31/40
A	EP-A-0 167 253 (THE UPJOHN CO.) * page 2, lines 1-3,10-15; page 3 * ---	1,13	
A	DE-A-2 749 950 (THE UPJOHN CO.) * pages 1-3, claim 1; page 16, claim 23 *; & US - A - 4 145 435 (Cat. D) ---	1,13	
A	DE-A-3 241 933 (THE UPJOHN CO.) * pages 1,2, claim 1; page 14, lines 15-22 *; & US - A - 4 438 130 (Cat. D) -----	1,13	
			TECHNICAL FIELDS SEARCHED (Int. Cl.5)
			C 07 D 295/00 C 07 D 405/00
The present search report has been drawn up for all claims			
Place of search BERLIN		Date of completion of the search 18-04-1990	Examiner KYRIAKAKOU G
CATEGORY OF CITED DOCUMENTS			
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